8.0 Summary of Safety and Effectiveness Data, Data Available Statement

I certify that, in my capacity as President of Medical Metrics, Inc., data demonstrating the safety and effectiveness of the KIMAX™ system is on file and is available upon request to any person within 30 days of a written request. The information will exclude all patient identifiers, and trade secret and confidential commercial information, as defined in 21 CFR 20.61.

Signature

Frank Vazquez, President, Medical Metrics, Inc.

Jan. 18, 2001
Date

K _____ (pending)
Premarket Notification Number

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

FEB 1 4 2001

Medical Metrics, Inc.
C/O Dr. Ann Tunstall
Salamandra, LLC
4600 North Park Ave.
Suite 100
CHEVY CHASE MD 20815-4518

Re: K010187
KIMAX, Model 1024 (Fluoroscopic Image System)
Dated: January 18, 2001
Received: January 22, 2001
Regulatory Class: II
21 CFR §892.1650/Procode: 90 JAA

Dear Dr. Tunstall:

We have reviewed your Section 510(k) notification of intent to market the device referenced above and we have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (Premarket Approval), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 895. A substantially equivalent determination assumes compliance with the Current Good Manufacturing Practice requirements, as set forth in the Quality System Regulation (QS) for Medical Devices: General regulation (21 CFR Part 820) and that, through periodic QS inspections, the Food and Drug Administration (FDA) will verify such assumptions. Failure to comply with the GMP regulation may result in regulatory action. In addition, FDA may publish further announcements concerning your device in the Federal Register. Please note: this response to your premarket notification submission does not affect any obligation you might have under sections 531 through 542 of the Act for devices under the Electronic Product Radiation Control provisions, or other Federal laws or regulations.

This letter will allow you to begin marketing your device as described in your 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801 and additionally 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4639. Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its internet address "http://www.fda.gov/cdrh/dsma/dsmamain.html".

Sincerely yours,

Daniel G. Schultz, M.D.
Captain, USPHS
Acting Director, Division of Reproductive,
Abdominal, and Radiological Devices
Office of Device Evaluation
Center for Devices and Radiological Health

Enclosure (s)

510(k) Number (if known): K010187

Device Name: KIMAX™ Model 1024

Indications For Use:

1.0 Statement of Intended Use

The KIMAX™ Model 1024 Kinematic Imaging and Musculoskeletal Assessment System is a diagnostic fluoroscopic X-ray imaging device. It is intended for fluoroscopic imaging of the musculoskeletal anatomy or motion between musculoskeletal anatomical structures.

The device can be used in most medical settings, including offices, clinics, outpatient imaging centers, mobile facilities and hospitals.

(PLEASE DO NOT WRITE BELOW THIS LINE - CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use ✓ OR Over-The-Counter Use_____
(Per 21 CFR 801.109)

(Optional Format 1-2-96)

David A. Seymour
(Division Sign-Off)
Division of Reproductive, Abdominal, ENT, and Radiological Devices
510(k) Number K010187